## What is claimed is:

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- A purification process for manufacturing a high pure acarbose uses alcohol for precipitation and separation, a strongly cation exchange chromatography and immobilized enzyme affinity chromatography for purification purifying acarbose-containing and an fermentation broth to get a high pure acarbose.
- The purification process of claim 1, wherein the strongly cation exchange chromatography uses a styrene
  divinylbenzene copolymer without methoxymethylmethacrylamide to be a resin matrix.
  - 3. The purification process of claim 1, wherein the enzyme of the immobilized enzyme affinity chromatography uses  $\alpha$  -amyloglucosidase( $\alpha$ -glucoamylase).
- 15 4. The purification process of claim 1, wherein the strongly cation exchange chromatography uses a cation exchange

resin containing 20-200 mg sugars/mL resin.

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- 5. The purification process of claim 2, wherein further comprising a step after the strongly cation exchange chromatography uses a solvent, 0~2.0N ammonia solution, to manufacture a high pure acarbose.
  - 6. The purification process as claim 3, wherein further comprising a step after the immobilized enzyme affinity chromatography uses a solvent, 55~75°C distilled water, to manufacture a high pure acarbose.
- 10 7. The purification process as claim 1, wherein the purity of high pure acarbose is large than 95% (wt/wt) used to treat diabetes.
  - 8. A purification process for purifying the acarbose comprising the steps of:
- 15 eliminating myselium from an acarbose-containing fermentation broth by centrifugation;

concentrating filtrate of the acarbose-containing fermentation broth to be consistency by a concenteration system;

adding adequate ethyl alcohol to the consistency and blending to be a solution;

taking an upper liquid from the solution by centrifugating; concentrating the upper liquid to be a consistency by the concentrating system;

putting the consistency into ethyl alcohol to get a 10 consistency liquid;

taking a sediment from the consistency liquid by centrifugating and solving the sediment by water to get an impure acarbose solution;

blending a strongly cation exchange resin with the 15 acarbose solution to get a resin;

using sodium chloride solution to eliminate an impurity in

## the resin;

using ammonia solution to eliminate an impurity in the resin; and solving the resin with ammonia solution to get a high pure

- 5 acarbose.
  - 9. The purification process as claim 8, wherein the eliminating myselium from acarbose-containing fermentation broth step could use a filter to replace centrifugating.
- 10 10. The purification process as claim 8, wherein the purity of high pure acarbose is 60%(wt/wt).
  - 11. A purification process for manufacturing a high pure acarbose comprising the steps of:
    - adjusting pH value of an impure acarbose;
- adding an cation exchange resin into the impure acarbose to get a solution;

blending the solution and taking an upper liquid;

adding a strong cation exchange resin into the upper liquid to get a mixing solution;

mixing and shaking the mixing solution to make the strong

5 cation exchange resin absorbing acarbose;

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using sodium chloride solution to eliminate an impurity in the acarbose; and

using ammonia solution to elute the acarbose to get a high pure acarbose.

- 10 12. The purification process as claim 12, wherein after the adjusting pH value step adds a cation exchange resin containing 250 mg sugars/g resin.
  - 13. The purification process as claim 12, wherein after taking the upper liquid adds a strong cation exchange resin containing 80 mg sugars/mL.
  - 14. The purification process as claim 12, wherein the purity of

high pure acarbose is up 78%.

15. A purification process for manufacturing a high pure acarbose comprising the steps of:

adjusting pH value of an upper liquid from an impure

5 acarbose mixing a strong cation exchange resin;

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passing the upper liquid through a strong cation exchange resin column;

washing the strong cation exchange resin in the column by deionized water till the absorbance of strong cation exchange resin being zero or steady;

getting an impure acarbose by using ammonia solution to elute the strong cation exchange resin;

concentrating the acarbose-containing fractions to be a volume by a concenteration system; and

using alcohol for extracting the impure acarbose to get a high pure acarbose.

- 16. The purification process as claim 16, wherein the flow velocity of passing the string cation exchange resin column is 2.5 mL/min.
- 17. The purification process of claim 16, wherein the ammonia solution gradient of ammonia solution for eluting the impure acarbose is 0.5~1.5N.
  - 18. The purification process as claim 16, wherein the purity of high pure acarbose is up 85%.
- 19. A purification process for manufacturing a high pure10 acarbose comprising the steps of:

solving a powder of acarbose, which the purity is 83%~87%, by distilled water to be a solution;

adjusting pH value of the solution;

passing the solution through  $\alpha$ -amyloglucosidase column;

washing the  $\alpha$ -amyloglucosidase column by using a times deionied water volume as the volume of the  $\alpha$ 

-amyloglucosidase column;

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eluting an acarbose from the lpha -amyloglucosidase column by distilled water;

concentrating the acarbose-containing fractions to be a volume by a concenteration system; and

using alcohol for precipitating the impure acarbose to get a high pure acarbose.

- 20. The purification process of claim 20, wherein the flow velocity of passing the  $\alpha$ -amyloglucosidase column is 1.5 mL/min.
- 21. The purification process of claim 20, wherein the washing the  $\alpha$  -amyloglucosidase column step uses two times deionized water volume as the volume of the  $\alpha$  -amyloglucosidase column.
- 15 22. The purification process of claim 20, wherein washing the  $\alpha$ -amyloglucosidase column by deionized water step

changes the flow velocity of passing the  $\alpha$  -amyloglucosidase column being 210nm till the absorbance of the  $\alpha$  -amyloglucosidase is steady.

- 23. The purification process of claim 20, wherein solving an impure acarbose from the  $\alpha$ -amyloglucosidase column by distilled water, 65°C.
  - 24. The purification process of claim 20, wherein the purity of the high pure acarbose is up 95%.